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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(54) Title: <b>GENE REPAIR INVOLVING <i>IN VIVO</i> EXCISION OF TARGETING DNA</b></p> <p>(57) Abstract</p> <p>Methods of modifying, repairing, attenuating and inactivating a gene or other chromosomal DNA in a cell through SCEI-induced recombination are disclosed. Also disclosed are methods of treating or prophylaxis of a genetic disease in an individual in need thereof.</p>			

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GENE REPAIR INVOLVING  
*IN VIVO* EXCISION OF TARGETING DNA

RELATED APPLICATION

This application claims the benefit of U.S. Provisional Application  
5 No. 60/118,472, filed February 3, 1999, the entire teachings of which are  
incorporated herein by reference.

BACKGROUND OF THE INVENTION

Homologous recombination and, more specifically D-loop mediated recombination, provide a method for genetically modifying chromosomal DNA sequences in a precise way. In addition to the possibility of introducing small precise mutations in order to alter the activity of the chromosomal DNA sequences, such a methodology makes it possible to correct the genetic defects in genes which can cause disease. Unfortunately, current methods for achieving homologous recombination are inherently inefficient, in that homologous recombination or D-loop recombination-mediated gene repair can usually be achieved in only a small proportion of cells that have taken up the relevant "targeting or correcting" DNA. For example, in cultured mammalian cells, such recombinational events usually occur in only one in ten thousand cells which have taken up the relevant targeting or correcting DNA. Accordingly, the use of biochemical selections are normally necessary to identify and isolate cells which have successfully recombined input DNA.

Thus, there is a need to develop new and improved methods of homologous recombination or D-loop recombination-mediated gene repair.

SUMMARY OF THE INVENTION

25 The present invention is related to Applicants' discovery that excision of targeting or correcting DNA from a vector within cells which have taken up the

vector significantly increased the frequency of homologous recombination and D-loop recombination-mediated gene repair in these cells. As a result, Applicants' invention relates to methods which result in excision of targeting or correcting DNA from a vector within cells which have taken up the vector. The methods comprise

5 introducing into a cell (a) a first vector which comprises a targeting DNA, wherein the targeting DNA comprises DNA homologous to a chromosomal target site and is flanked by specific restriction endonuclease site(s), and (b) a restriction endonuclease which cleaves the restriction endonuclease site(s) and is present in the first vector or a second (separate) vector which comprises a nucleic acid encoding

10 the restriction endonuclease or is introduced as the restriction endonuclease itself. In one embodiment, two vectors are introduced into cells: a first vector which comprises a targeting DNA, wherein the targeting DNA comprises DNA homologous to a chromosomal target site and is flanked by specific restriction endonuclease sites and a second vector which comprises a nucleic acid (e.g., DNA)

15 which encodes the restriction endonuclease. Alternatively, a single vector which comprises both targeting DNA, wherein the targeting DNA comprises DNA homologous to a chromosomal target site and is flanked by specific restriction endonuclease site(s), and a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s). is introduced into the cell. In the

20 embodiments described herein, the targeting DNA is flanked by a restriction endonuclease site if such a site is present at or near either or both ends of the targeting DNA. That is, there can be one restriction endonuclease site present at or near one end of the targeting DNA or there can be two such sites, one at or near each end of the targeting DNA. The restriction endonuclease site(s) are recognized

25 (cleaved) by the restriction endonuclease used in the method. As described below, the endonuclease used in the method is one whose activity does not lead to the death of cells in which it cleaves. One example of an endonuclease useful in the method is a meganuclease enzyme. Two (or more) different restriction endonucleases can be used in the present method.

The present invention relates to a method of repairing a specific sequence of interest in chromosomal DNA of a cell comprising introducing into the cell (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site or sites and comprises (1) DNA homologous to

5   chromosomal DNA adjacent to the specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. The two can be introduced, as described above, in the same or separate vectors or a vector comprising targeting

10   DNA flanked by specific restriction endonuclease site(s) and the endonuclease itself (not in a vector) can be introduced. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. Typically, the targeting DNA is designed such that the homologous DNA is at the left and right arms of the targeting DNA construct and DNA which repairs the specific sequence of interest is inserted between the two

15   arms. In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector which comprises a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding the restriction endonuclease

20   which cleaves the specific sites present in the vector are introduced into the cell in the same vector. As used herein, chromosomal DNA adjacent to a specific sequence of interest refers to chromosomal DNA present near or next to the specific sequence of interest.

In a particular embodiment, the specific sequence of interest is a mutation.

25   The present invention also relates to a method of modifying a specific sequence (or gene) in chromosomal DNA of a cell comprising introducing into the cell (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the specific sequence (or gene) to be modified and (2) DNA which results in

30   modification of the specific sequence (or gene) upon recombination between the

targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites (one at or near each end of the targeting DNA). Typically, the targeting DNA is designed such that the

5 homologous DNA is at the left and right arms of the targeting DNA construct and DNA which results in modification of the specific sequence (or gene) is inserted between the two arms. In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector (either RNA or DNA) which comprises a nucleic acid encoding the restriction

10 endonuclease. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

The invention further relates to a method of attenuating an endogenous gene of interest in a cell comprising introducing into the cell (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to a target site of the endogenous gene of interest and (2) DNA which attenuates the gene of interest upon recombination between the targeting DNA and the gene of interest, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector.

15 Preferably, the targeting DNA is flanked by two restriction endonuclease sites. Typically, the targeting DNA is designed such that the homologous DNA is at the left and right arms of the targeting DNA construct and DNA which attenuates the gene of interest is located between the two arms. In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into

20 the cell a second vector (either RNA or DNA) which comprises a nucleic acid encoding the restriction endonuclease. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

The present invention also relates to a method of introducing a mutation into

30 a target site of chromosomal DNA of a cell comprising introducing into the cell (a) a

first vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the target site and (2) the mutation to be introduced into the chromosomal DNA, and (b) a second vector (RNA or DNA) comprising a nucleic acid encoding a restriction

- 5     endonuclease which cleaves the restriction endonuclease site present in the first vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. Typically, the targeting DNA is designed such that the homologous DNA is at the left and right arms of the targeting DNA construct and the mutation is located between the two arms. In another embodiment of this method, the restriction
- 10    endonuclease is introduced directly into the cell. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site are introduced into the cell in the same vector.

The present invention also relates to the resulting cells and to their uses, such as for production of proteins or other gene products or for treatment or prophylaxis of a condition or disorder in an individual (e.g., a human or other mammal or vertebrate) arising as a result of a genetic defect (mutation). For example, cells can be produced (e.g., *ex vivo*) by the methods described herein and then introduced into an individual using known methods. Alternatively, cells can be modified in the

- 20    individual (without being removed from the individual).

Thus, the invention further relates to a method of treating or prophylaxis of a genetic disease in an individual in need thereof. In one embodiment, this method comprises introducing into the individual cells which comprise (a) a first vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site or sites and comprises (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) a second vector (RNA or DNA) comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the first vector. In a second embodiment, this method comprises

introducing into the individual cells which comprise (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site(s) and comprises (1) DNA homologous to chromosomal DNA and (2) DNA which repairs the specific sequence of interest upon recombination between the

5 targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. In a third embodiment, this method comprises introducing into the individual cells which comprise a vector comprising (a) targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site(s) and comprises (1) DNA homologous to

10 chromosomal DNA and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the plasmid. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. Typically, the targeting DNA is designed

15 such that the homologous DNA is at the left and right arms of the targeting DNA construct and DNA which repairs the specific sequence of interest is located between the two arms.

Alternatively, in a method of treating or prophylaxis of a genetic disease in an individual in need thereof, restriction endonucleases and vectors comprising

20 targeting DNA and/or nucleic acid encoding a restriction endonuclease can be administered directly to the individual.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic diagram of an embodiment of a homologous recombination or D-loop recombination-mediated repair method described herein.

25 Figure 2 is a table which provides the results from I-SceI induced D-loop recombination-mediated repair experiments in NIH3T3 cells.

Figure 3 is a table providing examples of meganuclease enzymes.

#### DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to the development of a generally useful method for significantly increasing the frequency of homologous recombination and D-loop recombination-mediated gene repair. At least *in vitro*, over 1% of a population of transfected cells can be shown to generate the desired recombinational events using the methods described herein. It is likely that these findings represent the ability to achieve homologous recombination and/or gene repair in close to 10% of successfully transfected cells (or higher) when corrected for the efficiency of transfection (the percent cells that take up DNA).

The invention relates to the use of methods which lead to the excision of homologous targeting DNA sequences from a recombinant vector within transfected cells (cells which have taken up the vector). The methods comprise introducing into cells (a) a first vector which comprises a targeting DNA, wherein the targeting DNA flanked by specific restriction endonuclease site(s) and comprises DNA homologous to a chromosomal target site, and (b) a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the first vector or a second vector which comprises a nucleic acid encoding the restriction endonuclease. Alternatively, a vector which comprises both targeting DNA and a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s) is introduced into the cell. Nucleic acid encoding the restriction endonuclease is also referred to herein as an expression cassette encoding the restriction endonuclease. Targeting DNA is also referred to herein as a repair matrix and correcting DNA.

In the embodiments described herein, the targeting DNA is flanked by a restriction endonuclease site if such a site is present at or near either or both ends of the targeting DNA. That is, there can be one restriction endonuclease site present at or near one end of the targeting DNA or there can be two such sites, one at or near each end of the targeting DNA.

A restriction endonuclease used in the present invention recognizes a target DNA sequence (e.g., a restriction endonuclease site) which would not lead to death of the cells upon cleavage of the DNA sequence by the restriction endonuclease. A meganuclease enzyme, which recognizes a very large DNA sequence, is an example

of a restriction endonuclease which can be used in the present invention. An example of a meganuclease enzyme is I-SceI which recognizes an 18-bp site (DNA sequence) that does not appear to be represented in murine or human DNA. Other examples of meganuclease enzymes are provided in Figure 3. Other meganuclease 5 enzymes (natural and synthetic) are known and described in the art. In a particular embodiment, a restriction endonuclease used in the present invention has a specificity of at least  $6.7 \times 10^{-10}$  of cleaving (cutting) frequency.

Expression of commonly used four and six base cutting restriction enzymes within cells would usually lead to cleavage of chromosomal DNA and death of the 10 cells due to the existence of many restriction sites within the cellular DNA which are recognized by the enzymes. Accordingly, such restriction enzymes are not used in the present invention.

The excision of a linear segment of DNA within cells (presumably within the nucleus) appears to generate a form of DNA which can be more efficiently utilized 15 for recombination than either circular DNA or DNA linearized *in vitro* (prior to transfection) that are introduced into cells. This may relate to the generation of a linear segment of DNA that is either more resistant to exonucleolytic degradation than linear DNA that is transfected, or perhaps to the generation of a template more capable of forming complexes with gene products essential for recombinational 20 event.

The ability to achieve homologous recombination and gene repair at high efficiency allows for the treatment of genetic diseases by true gene repair, rather than by the addition of a functional gene to genes, as is currently the major focus of gene therapy. The method described herein should not require long term expression 25 of introduced DNA *in vivo*, a common problem with current gene therapy experiments, since only the transient expression of the appropriate restriction endonuclease should be necessary to excise the 'correcting' linear segment of DNA.

The present invention relates to a method of repairing a specific sequence of interest in chromosomal DNA of a cell comprising introducing into the cell (a) a 30 vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site or sites and comprises (1) DNA homologous to

chromosomal DNA adjacent to the specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. Preferably, the targeting 5 DNA is flanked by two restriction endonuclease sites (one at or near each end of the targeting DNA). In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector which comprises a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. In yet another embodiment 10 of this method, both targeting DNA and nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

In a method of repairing a specific sequence of interest in chromosomal DNA of a cell, the targeting DNA is designed such that homologous recombination, and more preferably, D-loop mediated recombination, occurs between the targeting 15 DNA and chromosomal DNA and, upon recombination, repair of the specific sequence of interest occurs. Thus, in a particular embodiment, the targeting DNA is designed to include (1) DNA homologous to chromosomal DNA adjacent to the specific sequence of interest, wherein the homologous DNA is sufficient for recombination between the targeting DNA and chromosomal DNA, and (2) DNA 20 which repairs the specific sequence of interest upon recombination between the targeting DNA and chromosomal DNA. Typically, the homologous DNA of the targeting DNA construct flanks each end of the DNA which repairs the specific sequence of interest. That is, the homologous DNA is at the left and right arms of the targeting DNA construct and the DNA which repairs the sequence of interest is 25 located between the two arms.

In a particular embodiment, the specific sequence of interest is a mutation. Thus, in this embodiment, the invention relates to a method of repairing a mutation in chromosomal DNA of a cell comprising introducing into the cell (a) a vector comprising targeting DNA wherein the targeting DNA is flanked by a restriction 30 endonuclease site or sites and comprises (1) DNA homologous to chromosomal

DNA adjacent to the mutation and (2) DNA which repairs the mutation upon recombination between the targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites (one at or near each end of the targeting DNA). In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector which comprises a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site(s) present in the vector. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

In a method of repairing a mutation in chromosomal DNA of a cell, the targeting DNA is designed such that homologous recombination, and more preferably, D-loop mediated recombination, occurs between the targeting DNA and chromosomal DNA and, upon recombination, repair of the mutation occurs. Thus, in a particular embodiment, the targeting DNA is designed to include (1) DNA homologous to chromosomal DNA adjacent to the mutation, wherein the homologous DNA is sufficient for recombination between the targeting DNA and chromosomal DNA, and (2) DNA which repairs the mutation upon recombination between the targeting DNA and chromosomal DNA. Typically, the homologous DNA of the targeting DNA construct flanks each end of the DNA which repairs the mutation. That is, the homologous DNA is at the left and right arms of the targeting DNA construct and the DNA which repairs the mutation is located between the two arms.

As used herein, a mutation refers to a nucleotide change, such as a single or multiple nucleotide substitution, deletion or insertion, in a nucleotide sequence. Preferably, the mutation is a point mutation. Chromosomal DNA which bears a mutation has a nucleic acid sequence that is different in sequence from that of the corresponding wildtype chromosomal DNA.

As used herein, chromosomal DNA adjacent to a specific sequence of interest refers to chromosomal DNA present near or next to the specific sequence of interest.

The present invention also relates to a method of modifying a specific sequence (or gene) in chromosomal DNA of a cell comprising introducing into the cell (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the specific sequence (or gene) to be modified and (2) DNA which modifies the specific sequence (or gene) upon recombination between the targeting DNA and the 5 chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector (either RNA or DNA) which comprises a nucleic acid encoding the 10 restriction endonuclease. In yet another embodiment of this method, both targeting 15 DNA and nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

In a method of modifying a specific sequence (or gene) in chromosomal DNA of a cell, the targeting DNA is designed such that homologous recombination, 20 and more preferably, D-loop mediated recombination, occurs between the targeting DNA and chromosomal DNA and, upon recombination, modification of the sequence (or gene) occurs. Thus, in a particular embodiment, the targeting DNA is designed to include (1) DNA homologous to the specific sequence (or gene) to be modified, wherein the homologous DNA is sufficient for recombination between the 25 targeting DNA and chromosomal DNA, and (2) DNA which modifies the specific sequence (or gene) upon recombination between the targeting DNA and the chromosomal DNA. Typically, the homologous DNA of the targeting DNA construct flanks each end of the DNA which modifies the specific sequence (or gene). That is, the homologous DNA is at the left and right arms of the targeting

DNA construct and the DNA which modifies the specific sequence (or gene) is located between the two arms.

The invention further relates to a method of attenuating or inactivating an endogenous gene of interest in a cell comprising introducing into the cell (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to a target site of the endogenous gene of interest and (2) DNA which attenuates or inactivates the gene of interest upon recombination between the targeting DNA and the gene of interest, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites, as described above. In another embodiment of this method, the restriction endonuclease is introduced into the cell by introducing into the cell a second vector (either RNA or DNA) which comprises a nucleic acid encoding the restriction endonuclease. In yet another embodiment of this method, both the targeting DNA and the nucleic acid encoding the restriction endonuclease are introduced into the cell in the same vector.

In a method of attenuating or inactivating an endogenous gene of interest in a cell, the targeting DNA is designed such that homologous recombination, and more preferably, D-loop mediated recombination, occurs between the targeting DNA and endogenous gene of interest and, upon recombination, attenuation or inactivation of the gene of interest occurs. Thus, in a particular embodiment, the targeting DNA is designed to include (1) DNA homologous to a target site of the endogenous gene of interest, wherein the homologous DNA is sufficient for recombination between the targeting DNA and the gene of interest, and (2) DNA which attenuates or inactivates the gene of interest upon recombination between the targeting DNA and the gene of interest. Typically, the homologous DNA of the targeting DNA construct flanks each end of the DNA which attenuates or inactivates the gene of interest. That is, the homologous DNA is at the left and right arms of the targeting DNA construct and the DNA which attenuates or inactivates the gene of interest is located between the two arms.

The present invention also relates to a method of introducing a mutation into a target site (or gene) of chromosomal DNA of a cell comprising introducing into the cell (a) a first vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to 5 the target site (or gene) and (2) the mutation to be introduced into the chromosomal DNA, and (b) a second vector (RNA or DNA) comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. In another embodiment of this method, the restriction 10 endonuclease is introduced directly into the cell. In yet another embodiment of this method, both targeting DNA and nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site, are introduced into the cell in the same vector.

In a method of introducing a mutation into a target site (or gene) of 15 chromosomal DNA of a cell, the targeting DNA is designed such that homologous recombination, and more preferably, D-loop mediated recombination, occurs between the targeting DNA and the chromosomal DNA and, upon recombination, a mutation is introduced into the target site (or gene). Thus, in a particular embodiment, the targeting DNA is designed to include (1) DNA homologous to the 20 target site (or gene), wherein the homologous DNA is sufficient for recombination between the targeting DNA and the chromosomal DNA, and (2) the mutation which is introduced into the chromosomal DNA upon recombination between the targeting DNA and the chromosomal DNA. Typically, the homologous DNA of the targeting DNA construct flanks each end of the mutation. That is, the homologous DNA is at 25 the left and right arms of the targeting DNA construct and the mutation to be introduced into the chromosomal DNA (i.e., into a target site or gene) is located between the two arms.

The invention further relates to a method of treating or prophylaxis of a 30 genetic disease in an individual in need thereof. As used herein, a genetic disease refers to a disease or disorder that arises as a result of a genetic defect (mutation) in

a gene in the individual. In a particular embodiment, the genetic disease arises as a result of a point mutation in a gene in the individual.

In one embodiment, the method of treating or prophylaxis of a genetic disease in an individual in need thereof comprises introducing into (administering 5 to) the individual cells which comprise (a) a first vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) 10 a second vector (RNA or DNA) comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector. In a second embodiment, the method comprises introducing into the individual cells which comprise (a) a vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA 15 homologous to chromosomal DNA adjacent to a specific sequence of interest (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. In a third embodiment, the method comprises introducing into the individual cells which 20 comprise a vector comprising (a) targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) nucleic acid encoding a restriction 25 endonuclease which cleaves the restriction endonuclease site present in the plasmid. Preferably, the targeting DNA is flanked by two restriction endonuclease sites. Typically, the homologous DNA of the targeting DNA construct flanks each end of the DNA which repairs the specific sequence of interest. That is, the homologous DNA is at the left and right arms of the targeting DNA construct and the DNA 30 which repairs the sequence of interest is located between the two arms.

Alternatively, in a method of treating or prophylaxis of a genetic disease in an individual in need thereof, restriction endonucleases and vectors comprising targeting DNA and/or nucleic acid encoding a restriction endonuclease can be administered directly to the individual. The mode of administration is preferably at

5 the location of the target cells. In one embodiment, the method comprises introducing into (administering to) the individual (a) a first vector comprising targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon

10 recombination between the targeting DNA and the chromosomal DNA, and (b) a second vector (RNA or DNA) comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector. In a second embodiment, the method comprises introducing into the individual (a) a vector comprising targeting DNA, wherein the targeting DNA is

15 flanked by a restriction endonuclease site and (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) a restriction endonuclease which cleaves the restriction endonuclease site present in the vector. In a third embodiment, the method

20 comprises introducing into the individual a vector comprising (a) targeting DNA, wherein the targeting DNA is flanked by a restriction endonuclease site and (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and DNA which repairs the specific sequence of interest upon recombination between the targeting DNA and the chromosomal DNA, and (b) nucleic acid

25 encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the plasmid. Preferably, the targeting DNA is flanked by two restriction endonuclease sites.

The invention also relates to the generation of animal models of disease in which restriction endonuclease sites (e.g., I-SceI target sites) are introduced at the

30 site of the disease gene for evaluation of optimal delivery techniques.

The efficiency of gene modification/repair can be enhanced by the addition expression of other gene products. The restriction endonuclease and other gene products can be directly introduced into a cell in conjunction with the correcting DNA or via RNA expression. The approach is applicable to all organisms.

5 Targeting DNA can be manufactured according to methods generally known in the art. For example, targeting DNA can be manufactured by chemical synthesis or recombinant DNA/RNA technology (see, e.g., Sambrook *et al.*, Eds., *Molecular Cloning: A Laboratory Manual*, 2nd edition, Cold Spring Harbor University Press, New York (1989); and Ausubel *et al.*, Eds., *Current Protocols In Molecular*

10 *Biology*, John Wiley & Sons, New York (1997)).

A "target site", as used herein, refers to a distinct chromosomal location at which a chromosomal DNA sequence is to be modified in a precise way in accordance with the methods described herein.

15 As used herein, a "vector" includes a nucleic acid vector, e.g., a DNA vector, such as a plasmid, a RNA vector, virus or other suitable replicon (e.g., viral vector).

Viral vectors include retrovirus, adenovirus, parvovirus (e.g., adeno-associated viruses), coronavirus, negative strand RNA viruses such as orthomyxovirus (e.g., influenza virus), rhabdovirus (e.g., rabies and vesicular stomatitis virus), paramyxovirus (e.g. measles and Sendai), positive strand RNA viruses such as picornavirus and alphavirus, and double stranded DNA viruses including adenovirus, herpesvirus (e.g., Herpes Simplex virus types 1 and 2, Epstein-Barr virus, cytomegalovirus), and poxvirus (e.g., vaccinia, fowlpox and canarypox). Other viruses include Norwalk virus, togavirus, flavivirus, reoviruses, papovavirus, hepadnavirus, and hepatitis virus, for example. Examples of 20 retroviruses include: avian leukosis-sarcoma, mammalian C-type, B-type viruses, D-type viruses, HTLV-BLV group, lentivirus, spumavirus (Coffin, J.M., *Retroviridae: The viruses and their replication*, *In Fundamental Virology*, Third Edition, B.N. Fields, *et al.*, Eds., Lippincott-Raven Publishers, Philadelphia, 1996). Other examples include murine leukemia viruses, murine sarcoma viruses, mouse 25 mammary tumor virus, bovine leukemia virus, feline leukemia virus, feline sarcoma

virus, avian leukemia virus, human T-cell leukemia virus, baboon endogenous virus, Gibbon ape leukemia virus, Mason Pfizer monkey virus, simian immunodeficiency virus, simian sarcoma virus, Rous sarcoma virus and lentiviruses. Other examples of vectors are described, for example, in McVey et al., U.S. Patent No. 5,801,030, 5 the teachings of which are incorporated herein by reference.

A vector comprising a nucleic acid encoding a restriction endonuclease contains all or part of the coding sequence for the restriction endonuclease operably linked to one or more expression control sequences whereby the coding sequence is under the control of transcription signals to permit production or synthesis of the 10 restriction endonuclease. Such expression control sequences include promoter sequences, enhancers, and transcription binding sites. Selection of the promoter will generally depend upon the desired route for expressing the restriction endonuclease. The elements can be isolated from nature, modified from native sequences or manufactured *de novo* (e.g., by chemical synthesis or recombinant DNA/RNA 15 technology, according to methods known in the art (see, e.g., Sambrook et al., Eds., *Molecular Cloning: A Laboratory Manual*, 2nd edition, Cold Spring Harbor University Press, New York (1989); and Ausubel et al., Eds., *Current Protocols In Molecular Biology*, John Wiley & Sons, New York (1997)). The elements can then be isolated and fused together by methods known in the art, such as exploiting and 20 manufacturing compatible cloning or restriction sites.

Similarly, a vector comprising targeting DNA flanked by a restriction endonuclease site can be manufactured according to methods generally known in the art. For example, the vector comprising targeting DNA flanked by a restriction endonuclease site can be manufactured by chemical synthesis or recombinant 25 DNA/RNA technology (see, e.g., Sambrook et al., Eds., *Molecular Cloning, A Laboratory Manual*, 2nd edition, Cold Spring Harbor University Press, New York, 1989; and Ausubel et al., Eds., *Current Protocols In Molecular Biology*, John Wiley & Sons, New York, 1994-1997).

Vectors comprising targeting DNA flanked by a restriction endonuclease site 30 and/or nucleic acid encoding a restriction endonuclease can be introduced into a cell

by a variety of methods (e.g., transformation, transfection, direct uptake, projectile bombardment, using liposomes). Examples of suitable methods of transfecting or transforming cells include calcium phosphate precipitation, electroporation, microinjection, infection, lipofection and direct uptake. Such methods are described 5 in more detail, for example, in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*, Second Edition, Cold Spring Harbor University Press, New York (1989); and Ausubel, *et al.*, *Current Protocols in Molecular Biology*, John Wiley & Sons, New York (1998), the teachings of which are incorporated herein by reference.

A vector comprising targeting DNA flanked by a restriction endonuclease 10 site and/or nucleic acid encoding a restriction endonuclease can also be introduced into a cell by targeting the vector to cell membrane phospholipids. For example, targeting of a vector of the present invention can be accomplished by linking the vector molecule to a VSV-G protein, a viral protein with affinity for all cell membrane phospholipids. Such a construct can be produced using methods well 15 known to those practiced in the art.

Restriction endonucleases can be introduced into a cell according to methods generally known in the art which are appropriate for the particular restriction endonuclease and cell type. For example, a restriction endonuclease can be 20 introduced into a cell by direct uptake, microinjection, calcium phosphate precipitation, electroporation, infection, and lipofection. Such methods are described in more detail, for example, in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*, Second Edition, Cold Spring Harbor University Press, New York (1989); and Ausubel, *et al.*, *Current Protocols in Molecular Biology*, John Wiley & Sons, New York (1998). Other suitable methods are also described in the 25 art. The restriction endonuclease can be coupled to a facilitator of protein entry to facilitate introduction of the enzyme into a cell. Examples of facilitators of protein entry include tat, HSV VP22 and anthrax toxin. Coupling of a protein to a facilitator of protein entry can be accomplished using methods well known to those practiced in the art. Protein delivery strategies (e.g., HSV VP22, anthrax toxin) can be 30 evaluated in accordance with the methods of the invention described herein.

Once in the cell, the restriction endonuclease and the vector comprising targeting DNA flanked by a restriction endonuclease site and/or nucleic acid encoding a restriction endonuclease are imported or translocated by the cell from the cytoplasm to the site of action in the nucleus.

5 As used herein, a cell refers to a prokaryotic cell, such as a bacterial cell, or eukaryotic cell, such as an animal, plant or yeast cell. A cell which is of animal or plant origin can be a stem cell or somatic cell. Suitable animal cells can be of, for example, mammalian, avian or invertebrate origin. Examples of mammalian cells include human (such as HeLa cells), bovine, ovine, porcine, murine (such as 10 embryonic stem cells), rabbit and monkey (such as COS1 cells) cells. The cell may be an embryonic cell, bone marrow stem cell or other progenitor cell. Where the cell is a somatic cell, the cell can be, for example, an epithelial cell, fibroblast, smooth muscle cell, blood cell (including a hematopoietic cell, red blood cell, T-cell, B-cell, etc.), tumor cell, cardiac muscle cell, macrophage, dendritic cell, neuronal 15 cell (e.g., a glial cell or astrocyte), or pathogen-infected cell (e.g., those infected by bacteria, viruses, virusoids, parasites, or prions).

The cells can be obtained commercially or from a depository or obtained directly from an individual, such as by biopsy. The cells used can be obtained from an individual to whom they will be returned or from another/different individual of 20 the same or different species. For example, nonhuman cells, such as pig cells, can be modified to include a DNA construct and then introduced into a human. Such a treating procedure is sometimes referred to as *ex vivo* treatment. *Ex vivo* therapy has been described, for example, in Kasid *et al.*, *Proc. Natl. Acad. Sci. USA*, 87:473 (1990); Rosenberg *et al.*, *N. Engl. J. Med.*, 323:570 (1990); Williams *et al.*, *Nature*, 25 310:476 (1984); Dick *et al.*, *Cell*, 42:71 (1985); Keller *et al.*, *Nature*, 318:149 (1985); and Anderson *et al.*, United States Patent No. 5,399,346. Alternatively, the cells need not be isolated from the individual where, for example, it is desirable to deliver the vector to the individual in gene therapy.

As used herein, the term "individual" includes mammals, as well as other 30 animals (e.g., birds, fish, reptiles, insects). The terms "mammal" and "mammalian",

as used herein, refer to any vertebrate animal, including monotremes, marsupials and placental, that suckle their young and either give birth to living young (eutharian or placental mammals) or are egg-laying (metatharian or nonplacental mammals).

Examples of mammalian species include humans and other primates (e.g., monkeys, 5 chimpanzees), rodents (e.g., rats, mice, guinea pigs) and ruminents (e.g., cows, pigs, horses).

Restriction endonucleases and vectors which comprise targeting DNA flanked by a restriction endonuclease site and/or nucleic acid encoding a restriction endonuclease can be introduced into an individual using routes of administration 10 generally known in the art (e.g., parenteral, mucosal, nasal, injection, systemic, implant, intraperitoneal, oral, intradermal, transdermal (e.g., in slow release polymers), intramuscular, intravenous including infusion and/or bolus injection, subcutaneous, topical, epidural, buccal, rectal, vaginal, etc.). The restriction endonucleases and vectors can, preferably, be administered in a pharmaceutically 15 acceptable carrier, such as saline, sterile water, Ringer's solution, and isotonic sodium chloride solution. The mode of administration is preferably at the location of the target cells.

The dosage of restriction endonuclease or vector of the present invention administered to an individual, including frequency of administration, will vary 20 depending upon a variety of factors, including mode and route of administration; size, age, sex, health, body weight and diet of the recipient; nature and extent of symptoms of the disease or disorder being treated; kind of concurrent treatment, frequency of treatment, and the effect desired.

The present invention will now be illustrated by the following examples, 25 which are not to be considered limiting in any way.

## EXAMPLES

## Example 1 Plasmid Construction

All DNA manipulations used standard techniques and procedures. Such methods are described, for example, in Sambrook *et al.*, *Molecular Cloning: A 5 Laboratory Manual*, Second Edition, Cold Spring Harbor University Press, New York (1989); and Ausubel, *et al.*, *Current Protocols in Molecular Biology*, John Wiley & Sons, New York (1998). All synthetic oligonucleotides were synthesized on automated instruments using standard techniques.

The p2Wlac plasmid was constructed as follows: First, the pPytknlslacZ 10 plasmid (Henry *et al.*, *C. R. Acad. Sci. III*, 322(12):1061-1070 (1999)) was digested with the *SpeI* and *Hind*III restriction enzymes, resulting in excision from the plasmid of a 578 bp fragment containing the ATG start codon and 178 bp at the 5' end of the coding region of the *nslacZ* gene. Second, the oligonucleotide 15 5'-CTAGATGCATAGGGATAACAGGGTAAT-3' (SEQ ID NO:1), paired with 5'-AGCTATTACCCTGTTATCCCTATGCAT-3' (SEQ ID NO:2), was inserted into the *SpeI*-*Hind* III restriction sites of the pPytknlslacZ plasmid (Henry *et al.*, *C. R. Acad. Sci. III*, 322(12):1061-1070 (1999)) to produce the pWnlslacZ plasmid. Insertion of the oligonucleotide at the *SpeI*-*Hind* III restriction sites resulted in 20 destruction of the *SpeI* and *Hind*III restriction sites and insertion of a *Nsi*I restriction site and an *I-Sce*I restriction site. The pWnlslacZ plasmid was then digested with the *Nhe*I and *Bgl*II restriction enzymes, resulting in excision from the plasmid of a 0.6 kb fragment containing the stop codon and SV40 polyadenylation signal at the 3' end of the *nslacZ* gene. The oligonucleotide 25 5'-GATCATGCATAGGGATAACAGGGTAAT-3' (SEQ ID NO:3), paired with 5'-CTAGATTACCCTGTTATCCCTATGCAT-3' (SEQ ID NO:4), was inserted into the *Nhe*I-*Bgl*II restriction sites of the pWnlslacZ plasmid. Insertion of the oligonucleotide at the *Nhe*I-*Bgl*II restriction sites resulted in destruction of the *Nhe*I and the *Bgl*II restriction sites and insertion of an *I-Sce*I restriction site and a *Nsi*I restriction site. The result of these insertions is the p2Wlac plasmid in which the

*nslacZ* gene with the ATG start codon, 178 bp at the 5' end, stop codon and SV40 polyadenylation signal deleted, is inserted between two I-SceI sites. As a result of the deletion of the start codon and 178 bp at the 5' end of the coding region, *nslacZ* gene expression is inactivated.

5        The pWlac plasmid was constructed as follows: First, the pPytknlslacZ plasmid was digested with the *SpeI* and *Hind*III restriction enzymes, resulting in excision from the plasmid of a 578 bp fragment containing the ATG start codon and 178 bp at the 5' end of the coding region of the *nslacZ* gene. Second, the oligonucleotide 5'-CTAGATGCATAGGGATAACAGGGTAAT-3' (SEQ ID NO:1),  
10      paired with 5'-AGCTATTACCCCTGTTATCCCTATGCAT-3' (SEQ ID NO:2), was inserted into the *SpeI*-*Hind*III restriction sites of the pPytknlslacZ plasmid to produce the pWnlslacZ plasmid. Insertion at this restriction site resulted in destruction of the *SpeI* and *Hind*III restriction sites and the insertion of an *Nsi*I restriction site and an I-SceI restriction site. The pWnlslacZ plasmid was digested  
15      with the *Nhe*I and *Bgl*II restriction enzymes, resulting in excision from the plasmid of the 0.6 kb fragment containing the stop codon and SV40 polyadenylation signal at the 3' end of the *nslacZ* gene. The 5' extensions of the *Nhe*I-*Bgl*II restriction sites of the pWnlslacZ plasmid were converted to blunt ends by a filling-in reaction using T4 DNA polymerase. The blunted ends were then ligated together. The result is the  
20      pWlac plasmid in which the *nslacZ* gene with the ATG start codon, 178 bp at the 5' end, stop codon and SV40 polyadenylation signal deleted, is bounded at the 5' end by one I-SceI site; the 3' end of the *nslacZ* gene is not bounded by a I-SceI site. As a result of the deletion of the start codon and 178 bp at the 5' end of the coding region, *nslacZ* gene expression is inactivated.

25        The p-lac plasmid was constructed as follows: First, the pPytknlslacZ plasmid was digested with the *SpeI* and *Hind*III restriction enzymes, resulting in excision from the plasmid of a 578 bp fragment containing the ATG start codon and 178 bp at the 5' end of the coding region of the *nslacZ* gene. The 5' extensions of the *SpeI*-*Hind*III restriction sites of the pPytknlslacZ plasmid were converted to  
30      blunt ends by a filling-in reaction using T4 DNA polymerase. The blunted ends

were then ligated together to produce the p-lacZ plasmid. The p-lacZ plasmid was digested with the *Nhe*I and *Bgl*II restriction enzymes, resulting in excision from the plasmid of the 0.6 kb fragment containing the stop codon and SV40 polyadenylation signal at the 3' end of the *nslacZ* gene. The 5' extensions of the *Nhe*I-*Bgl*II 5 restriction sites of the pWnlslacZ plasmid were converted to blunt ends by a filling-in reaction using T4 DNA polymerase. The blunted ends were then ligated together. The result is the p-lac plasmid in which the *nslacZ* gene with the ATG start codon, 178 bp at the 5' end, stop codon and SV40 polyadenylation signal deleted, is not bounded at the 5' or 3' end by a *I-Sce*I site. As a result of the deletion of the start 10 codon and 178 bp at the 5' end of the coding region, *nslacZ* gene expression is inactivated.

The 2.8 kb linear fragment of the *nslacZ* gene used in the experiments described herein was obtained as follows: The pPytknlslacZ plasmid was digested with *Nhe*I and *Hind*III and a 2.8 kb fragment was purified by agarose gel 15 electrophoresis. This 2.8 kb fragment, referred to herein as the lac fragment, contains a fragment of the *nslacZ* gene with the ATG start codon, 178 bp at the 5' end, stop codon and SV40 polyadenylation signal deleted.

The pCMV *I-Sce*I(+) and pCMV *I-Sce*I(-) plasmids were described in Choulika *et al.*, *C. R. Acad. Sci. III*, 317(11):1013-1019 (1994).

20 The target plasmid pPytknlslacZDBcl was produced by digesting the pPytknlslacZ plasmid with the *Bcl*I restriction enzyme after demethylation of the plasmid. The 5' protruding ends were filled-in by the Klenow fragment of *E. coli* DNA polymerase I and religated. The result is insertion of a 4 base pair direct repeat in the sequence of the *nslacZ* gene resulting in a frame shift of the open 25 reading frame, thereby inactivating expression of the gene. Thus, the plasmid does not express the  $\beta$ -galactosidase protein.

The target plasmid pPytknlslacZ $\Delta$ Bcl was produced by digesting the pPytknlslacZ plasmid with the *Bcl*I restriction enzyme after demethylation of the plasmid. The 4 base pair 5' protruding ends were degraded by T4 DNA polymerase 30 and the resulting blunted ends religated. The result is deletion of 4 base pairs within

the sequence of the *nslacZ* gene resulting in a frame shift of the open reading frame, thereby inactivating expression of the gene. Thus, the plasmid does not the  $\beta$ -galactosidase protein.

The pUSVneo plasmid was described in Choulika *et al.*, *J. Virol.*,  
5 70(3):1792-1798 (1996).

**Example 2 Cell Line Production and D-loop Recombination: Correction Of A 4  
Base Pair Insertion**

5  $\mu$ g of the pPytknslacZDBcl plasmid and 5  $\mu$ g of the pUSVneo plasmid  
were co-transfected into  $5 \times 10^4$  NIH 3T3 cells (American Type Culture Collection)  
10 in a 35 mm petri dish (Falcon) using the CaPO<sub>4</sub> precipitation method. 48 hours after  
transfection, the tissue culture medium was supplemented with 600  $\mu$ g/ml of  
Geneticin (Gibco BRL). Antibiotic selection was maintained during selection of  
Geneticin resistant clones and during subcloning. Forty-eight (48) Geneticin  
resistant clones were isolated and grown independently in Dulbeccos modified  
15 Eagles Medium (DMEM), 10% calf serum, for 15 days before evaluating for the  
presence of the *nslacZ* gene.

To evaluate for presence of the *nslacZ* gene, DNA was extracted from cells  
in all 48 cultures of Geneticin resistant clones. Fragments of the *nslacZ* gene were  
amplified by polymerase chain reaction (PCR) as described in *BioFeedback in  
20 BioTechniques*, Hanley & J. P. Merlie, Vol. 10, No. 1, p. 56T (1991). Forty-six (46)  
of 48 clones were positive for the presence of the *nslacZ* gene.

Twenty-four (24) of the 46 clones positive for the presence of the *nslacZ*  
gene were evaluated for expression of the mutated *nslacZ* gene. To evaluate for  
expression of the mutated *nslacZ* gene, RNA was extracted from cells in the  
25 corresponding 24 cultures of Geneticin resistant clones. RNA encoding the mutated  
*nslacZ* gene was amplified by reverse transcriptase polymerase chain reaction (RT-  
PCR). The oligonucleotide primer 5'-TACACGCGTCGTGATTAGCGCCG-3'  
(SEQ ID NO:5) was used for lacZ reverse transcription. PCR was performed as

described in *BioFeedback in BioTechniques*, Hanley & J. P. Merlie, Vol. 10, No. 1, p. 56T (1991). Eleven (11) of 24 clones showed a positive reaction.

Southern blot analysis of the genomic DNA of these 11 clones was performed and 3 clones were shown to have less than 3 intact copies of the

5 pPytknlslacZDBcl construct.

Histochemical analysis of these 3 clones was performed by X-Gal staining as described in Bonnerot *et al.*, *Methods in Enzymology, Guide To Techniques In Mouse Development*, Academic Press, pp. 451-469 (1993). Two (2) of 3 clones showed expression of  $\beta$ -galactosidase in less than  $1 \times 10^6$  cells.  $\beta$ -galactosidase in

10 these cells is probably the result of intragenic recombination of the 4 bp direct repeat inserted into the *BcI* restriction site. Northern blot analysis of the mRNA expressed by the integrated pPytknlslacZDBcl construct showed very little expression for one of the clones (the one with no background expression) and strong signals for two other clones (the ones expressing  $\beta$ -galactosidase in less than  $1 \times 10^6$  cells). These 15 two cell lines, NIH 3T3 DBcl1 and NIH 3T3 DBcl2, were selected to be the targets to the D-loop recombination.

#### *Ex vivo Recombination In NIH 3T3 DBcl1 And NIH 3T3 DBcl2 Cell Lines*

Three sets of experiments were performed, in triplicate, using the NIH 3T3 DBcl1 and NIH 3T3 DBcl2 cell lines. Each set of experiment, in triplicate,

20 comprises 8 different cotransfections of DNA mixes as shown in Table 1. Transfections were performed in either  $5 \times 10^4$  NIH 3T3 DBcl1 cells or  $5 \times 10^4$  NIH 3T3 DBcl2 cells in a 60 mm petri dish (Falcon) using the CaPO<sub>4</sub> precipitation method.

TABLE 1

Mix Number	Expression Plasmid	Quantity	Repair Matrix	Quantity
5	pCMV I-SceI(+)	9 µg	p2Wlac	1 µg
	pCMV I-SceI(+)	9 µg	pWlac	1 µg
	pCMV I-SceI(+)	9 µg	p-lac	1 µg
	pCMV I-SceI(+)	9 µg	lac	1 µg
10	pCMV I-SceI(-)	9 µg	p2Wlac	1 µg
	pCMV I-SceI(-)	9 µg	pWlac	1 µg
	pCMV I-SceI(-)	9 µg	p-lac	1 µg
	pCMV I-SceI(-)	9 µg	lac	1 µg

96 hours after transfection, cells were stained for  $\beta$ -galactosidase expression in X-Gal and blue colony forming units (bcfu) were counted. The number of bcfu is the result of the D-loop correction in each of the experiment. Results are shown in 15 Figure 3.

Transfection of NIH 3T3 DBcl2 cells with mix number 1 (pCMV I-SceI(+), 9 µg; p2Wlac, 1 µg) gave a 3 to 5% of  $\beta$ -galactosidase positive clones (out of three experiments) as the higher rate of D-loop correction of the pPytknlslacZDBcl mutated plasmid. Thus, after transfection of  $1 \times 10^5$  cells with mix number 1, 96 20 individual cells were cloned by limit dilution according to standard methods: Cells were grown in DMEM, 10% calf serum, and analyzed for  $\beta$ -galactosidase expression. Five (5) of 71 clones showed more than  $1 \times 10^6$  cells expressing  $\beta$ -galactosidase (ranging between 5 to 80% of the cells). Southern blot analysis of 25 these 5 clones showed that 100% of the cells had their *nslacZ* gene with a *BclI* site recovered. The lack of correspondence between the expression of the intact *nslacZ* open reading frame and the total repair of the genome is probably the result of transgene variegation.

Example 3 Cell Line Production and D-loop Recombination: Correction Of A 4  
Base Pair Deletion

5 µg of the pPytknlslacZΔBcl plasmid and 5 µg of the pUSVneo plasmid were cotransfected in 5x10<sup>4</sup> NIH 3T3 cells (American Type Culture Collection) in a 5 35 mm petri dish (Falcon) using the CaPO<sub>4</sub> precipitation method. 48 hours after transfection, the tissue culture medium was supplemented with 600 µg/ml of Geneticin (Gibco BRL). Antibiotic selection was maintained during selection of Geneticin resistant clones and during subcloning. Forty-eight (48) Geneticin resistant clones were isolated and grown independently in Dulbeccos modified 10 Eagles Medium (DMEM), 10% calf serum, for 15 days before evaluating for the presence of the *nslacZ* gene.

To evaluate for presence of the *nslacZ* gene, DNA was extracted from cells in all 48 cultures of Geneticin resistant clones. Fragments of the *nslacZ* gene were amplified by PCR as described in *BioFeedback in BioTechniques*, Hanley & J. P. 15 Merlie, Vol. 10, No. 1, p. 56T (1991). Forty-eight (48) of 48 clones were positive for the presence of the *nslacZ* gene.

Twenty-four (24) of the 48 clones positive for the presence of the *nslacZ* gene were evaluated for expression of the mutated *nslacZ* gene. To evaluate for expression of the mutated *nslacZ* gene, RNA was extracted from cells in the 20 corresponding 24 cultures of Geneticin resistant clones. RNA encoding the mutated *nslacZ* gene was amplified by RT-PCR. The oligonucleotide primer 5'-TACACCGCGTCGTGATTAGCGCCG-3' (SEQ ID NO:5) was used for lacZ reverse transcription. PCR was performed as described in *BioFeedback in BioTechniques*, Hanley & J. P. Merlie, Vol. 10, No. 1, p. 56T (1991). Nine (9) of 24 25 clones showed a positive reaction.

Southern blot analysis of the genomic DNA of these 9 clones was performed and 1 clone was shown to have less than 3 intact copies of the pPytknlslacZΔBcl construct.

Histochemical analysis of these 4 clones was performed by X-Gal staining as 30 described in Bonnerot *et al.*, *Methods in Enzymology, Guide To Techniques In Mouse Development*, Academic Press, pp. 451-469 (1993). No clones showed

expression of  $\beta$ -galactosidase. No intragenic recombination can occur in these cell lines. Northern blot analysis of the mRNA expressed by the integrated pPytknlslacZ $\Delta$ Bcl construct showed very little expression for two of the clones and strong signals for the other two clones. These two cell lines, NIH 3T3  $\Delta$ Bcl1 and 5 NIH 3T3  $\Delta$ Bcl2, were selected to be the targets to the D-loop recombination.

*Ex vivo Recombination In NIH 3T3  $\Delta$ Bcl1 And NIH 3T3  $\Delta$ Bcl2 Cell Lines*

Three sets of experiments were performed, in triplicate, using the NIH 3T3  $\Delta$ Bcl1 and NIH 3T3  $\Delta$ Bcl2 cell lines. Each set of experiment, in triplicate, comprises 8 different cotransfections of DNA mixes as shown in Table 2.

10 Transfections were performed in either  $5 \times 10^4$  NIH 3T3  $\Delta$ Bcl1 cells or  $5 \times 10^4$  NIH 3T3  $\Delta$ Bcl2 cells in a 60 mm petri dish (Falcon) by the CaPO<sub>4</sub> precipitation method.

TABLE 2

Mix Number	Expression Plasmid	Quantity	Repair Matrix	Quantity
15	pCMV I-SceI(+)	9 $\mu$ g	p2Wlac	1 $\mu$ g
	pCMV I-SceI(+)	9 $\mu$ g	pWlac	1 $\mu$ g
	pCMV I-SceI(+)	9 $\mu$ g	p-lac	1 $\mu$ g
	pCMV I-SceI(+)	9 $\mu$ g	lac	1 $\mu$ g
20	pCMV I-SceI(-)	9 $\mu$ g	p2Wlac	1 $\mu$ g
	pCMV I-SceI(-)	9 $\mu$ g	pWlac	1 $\mu$ g
	pCMV I-SceI(-)	9 $\mu$ g	p-lac	1 $\mu$ g
	pCMV I-SceI(-)	9 $\mu$ g	lac	1 $\mu$ g

96 hours after transfection, cells were stained for  $\beta$ -galactosidase expression in X-Gal and blue colony forming units (bcfu) were counted. The number of bcfu is 25 the result of the D-loop correction in each of the experiment. Results are shown in Figure 3.

Transfection of NIH 3T3  $\Delta$ Bcl12 with mix number 1 (pCMV I-SceI(+), 9  $\mu$ g; p2Wlac, 1  $\mu$ g) gave a 1 to 3% of  $\beta$ -galactosidase positive clones (out of the three experiments) as the higher rate of D-loop correction of the pPytknlslacZ $\Delta$ Bcl mutated plasmid. Thus, after transfection of  $1 \times 10^5$  cells with mix number 1, 96

5 individual cells were cloned by limit dilution. Cells were grown in DMEM, 10% calf serum, and analyzed for  $\beta$ -galactosidase expression. Two (2) of 66 clones showed cells expressing  $\beta$ -galactosidase (ranging between 30 to 80% of the cells). Southern blot analysis of these 2 clones showed that 100% of the cells had their

10 nlslacZ gene with a Bcl I site recovered. The lack of correspondence between the expression of the intact nlslacZ open reading frame and the total repair of the genome is probably the result of transgene variegation.

**Example 4 I-SceI induced D-loop Recombination**

The pPytknlslacZD-Bcl construct is integrated into the genomic DNA of NIH 3T3 cells as described in Example 2. In these cells, the nlslacZDBcl gene is

15 transcribed but  $\beta$ -galactosidase expression is not detected ( $\beta$ -gal $^-$  cells).  $\beta$ -gal $^-$  cells are cotransfected with the p2Wlac plasmid containing two I-SceI sites and an expression vector coding for I-SceI endonuclease. The p2Wlac plasmid is linearized *in vivo* by the I-SceI endonuclease and correct the DBcl mutation by D-loop recombination. As a result, these cells contain a pPytknlslacZ plasmid that expresses

20  $\beta$ -galactosidase ( $\beta$ -gal $^+$  cells). A schematic diagram of this experiment is depicted in Figure 1.

The teachings of all the articles, patents and patent applications cited herein are incorporated by reference in their entirety.

While this invention has been particularly shown and described with

25 references to preferred embodiments thereof, it will be understood by those skilled in the art that various changes in form and details may be made therein without departing from the spirit and scope of the invention as defined by the appended claims.

## CLAIMS

What is claimed is:

1. A method of repairing a specific sequence of interest in chromosomal DNA of a cell comprising the steps of:
  - 5 a) introducing into said cell a first vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to the specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between said targeting DNA and the chromosomal DNA; and
  - b) introducing into said cell a second vector comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector.
2. The method of Claim 1 wherein the first vector is a viral vector.
- 15 3. The method of Claim 1 wherein the second vector is a viral vector.
4. The method of Claim 1 wherein the first vector is a plasmid.
5. The method of Claims 1 wherein said targeting DNA is flanked by two restriction endonuclease sites, one site present at or near the 5' end of said targeting DNA and one site present at or near the 3' end of said targeting
- 20 DNA.
6. The method of Claims 1 wherein the specific sequence of interest is a mutation.

7. A method of repairing a specific sequence of interest in chromosomal DNA of a cell comprising the steps of:
  - a) introducing into said cell a vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to the specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between said targeting DNA and the chromosomal DNA; and
  - b) introducing into the cell a restriction endonuclease which cleaves the restriction endonuclease site present in the vector.
8. The method of Claim 7 wherein the vector is a viral vector.
9. The method of Claim 7 wherein said targeting DNA is flanked by two restriction endonuclease sites.
10. The method of Claims 7 wherein the specific sequence of interest is a mutation.
11. A method of repairing a specific sequence of interest in chromosomal DNA of a cell comprising introducing into the cell a vector comprising (a) targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to the specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between said targeting DNA and the chromosomal DNA; and (b) a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site.
12. The method of Claim 11 wherein the vector is a viral vector.

13. The method of Claims 11 wherein said targeting DNA is flanked by two restriction endonuclease sites.
14. The method of Claims 11 wherein the specific sequence of interest is a mutation.
- 5 15. A method of modifying a specific sequence in chromosomal DNA of a cell comprising the steps of:
  - a) introducing into the cell a first vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the specific sequence to be modified and (2) DNA which modifies the specific sequence upon recombination between said targeting DNA and the chromosomal DNA; and
  - b) introducing into the cell a second vector comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector.
16. The method of Claim 15 wherein the first vector is a viral vector.
17. The method of Claim 15 wherein the second vector is a viral vector.
18. The method of Claim 17 wherein the first vector is a plasmid.
19. The method of Claims 15 wherein said targeting DNA is flanked by two restriction endonuclease sites.
20. A method of modifying a specific sequence in chromosomal DNA of a cell comprising the steps of:

- 10 a) introducing into the cell a vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the specific sequence to be modified and (2) DNA modifies the specific sequence upon recombination between said targeting DNA and the chromosomal DNA; and

15 b) introducing into the cell a restriction endonuclease which cleaves the restriction endonuclease site present in the vector, under conditions appropriate for the restriction endonuclease to cleave the restriction endonuclease site in the vector of step a).
21. The method of Claim 20 wherein the vector is a viral vector.
22. The method of Claim 20 wherein said targeting DNA is flanked by two restriction endonuclease sites.
23. A method of modifying a specific sequence in chromosomal DNA of a cell comprising introducing into the cell a vector comprising (a) targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the specific sequence to be modified and (2) DNA which results in modification of the specific sequence upon recombination between said targeting DNA and the chromosomal DNA; and (b) a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site.
24. The method of Claim 23 wherein the vector is a viral vector.
25. The method of Claims 23, wherein said targeting DNA is flanked by two restriction endonuclease sites.

26. A method of treating or prophylaxis of a genetic disease in an individual in need thereof comprising the steps of:

5           a) introducing into the individual cells which comprise a first vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon recombination between said targeting DNA and the chromosomal DNA; and

10           b) introducing into the individual a second vector comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector.

27. The method of Claim 26 wherein the first vector is a viral vector.

28. The method of Claim 27 wherein the second vector is a viral vector.

15           29. The method of Claim 27 wherein the first vector is a plasmid.

30. The method of Claims 24 wherein said targeting DNA is flanked by two restriction endonuclease sites.

31. A method for treating or prophylaxis of a genetic disease in an individual in need thereof comprising the steps of:

20           a) introducing into the individual cells which comprise a vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to chromosomal DNA adjacent to a specific sequence of interest and (2) DNA which repairs the specific sequence of interest upon

recombination between said targeting DNA and the chromosomal DNA; and

b) introducing into the cell a restriction endonuclease which cleaves the restriction endonuclease site present in the vector.

5 32. The method of Claim 31 wherein the vector is a viral vector.

33. The method of Claim 31 wherein said targeting DNA is flanked by two restriction endonuclease sites.

34. A method of attenuating an endogenous gene of interest in a cell comprising the steps of:

10 a) introducing into the cell a first vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to a targeting site of the endogenous gene of interest and (2) DNA which attenuates the gene of interest upon recombination between said targeting DNA and the gene of interest; and

15 b) introducing into the cell a second vector comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector.

35. The method of Claim 34 wherein the first vector is a viral vector.

20 36. The method of Claim 34 wherein the second vector is a viral vector..

37. The method of Claim 34 wherein the first vector is a plasmid.

38. The method of Claims 34 wherein said targeting DNA is flanked by two restriction endonuclease sites.

39. A method of attenuating an endogenous gene of interest in a cell comprising the steps of:

- a) introducing into the cell a vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to a targeting site of the endogenous gene of interest and (2) DNA which attenuates the gene of interest upon recombination between said targeting DNA and the gene of interest; and
- b) introducing into the cell a restriction endonuclease which cleaves the restriction endonuclease site present in the vector.

40. The method of Claim 39 wherein the vector is a viral vector.

41. The method of Claim 39 wherein said targeting DNA is flanked by two restriction endonuclease sites.

42. A method of attenuating an endogenous gene of interest in a cell comprising introducing into the cell a vector comprising (a) targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to a targeting site of the endogenous gene of interest and (2) DNA which attenuates the gene of interest upon recombination between said targeting DNA and the gene of interest; and (b) a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site.

43. The method of Claim 42 wherein the vector is a viral vector.

44. The method of Claims 42 wherein said targeting DNA is flanked by two restriction endonuclease sites.

45. A method of introducing a mutation into a targeting site of chromosomal DNA of a cell comprising the steps of:

- a) introducing into the cell a first vector comprising targeting DNA, wherein said is flanked by a restriction endonuclease site and comprises (1) DNA homologous to said target site and (2) the mutation to be introduced into the chromosomal DNA; and
- b) introducing into the cell a second vector comprising a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site present in the first vector.

10 46. The method of Claim 45 wherein the first vector is a viral vector.

47. The method of Claim 45 wherein the second vector is a viral vector.

48. The method of Claim 47 wherein the first vector is a plasmid.

49. The method of Claims 45 wherein said targeting DNA is flanked by two restriction endonuclease sites.

15 50. A method of introducing a mutation into a target site of chromosomal DNA of a cell comprising the steps of:

- a) introducing into the cell a vector comprising targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the target site and (2) the mutation to be introduced into the chromosomal DNA; and
- b) introducing into the cell a restriction endonuclease which cleaves the restriction endonuclease site present in the vector.

20 51. The method of Claim 50 wherein the vector is a viral vector.

52. The method of Claim 50 wherein said targeting DNA is flanked by two restriction endonuclease sites.
53. A method of introducing a mutation into a target site of chromosomal DNA of a cell comprising introducing into the cell a vector comprising (a) targeting DNA, wherein said targeting DNA is flanked by a restriction endonuclease site and comprises (1) DNA homologous to the target site and (2) the mutation to be introduced into the chromosomal DNA; and (b) a nucleic acid encoding a restriction endonuclease which cleaves the restriction endonuclease site.
- 10 54. The method of Claim 53 wherein the vector is a viral vector.
55. The method of Claims 53 wherein said targeting DNA is flanked by two restriction endonuclease sites.

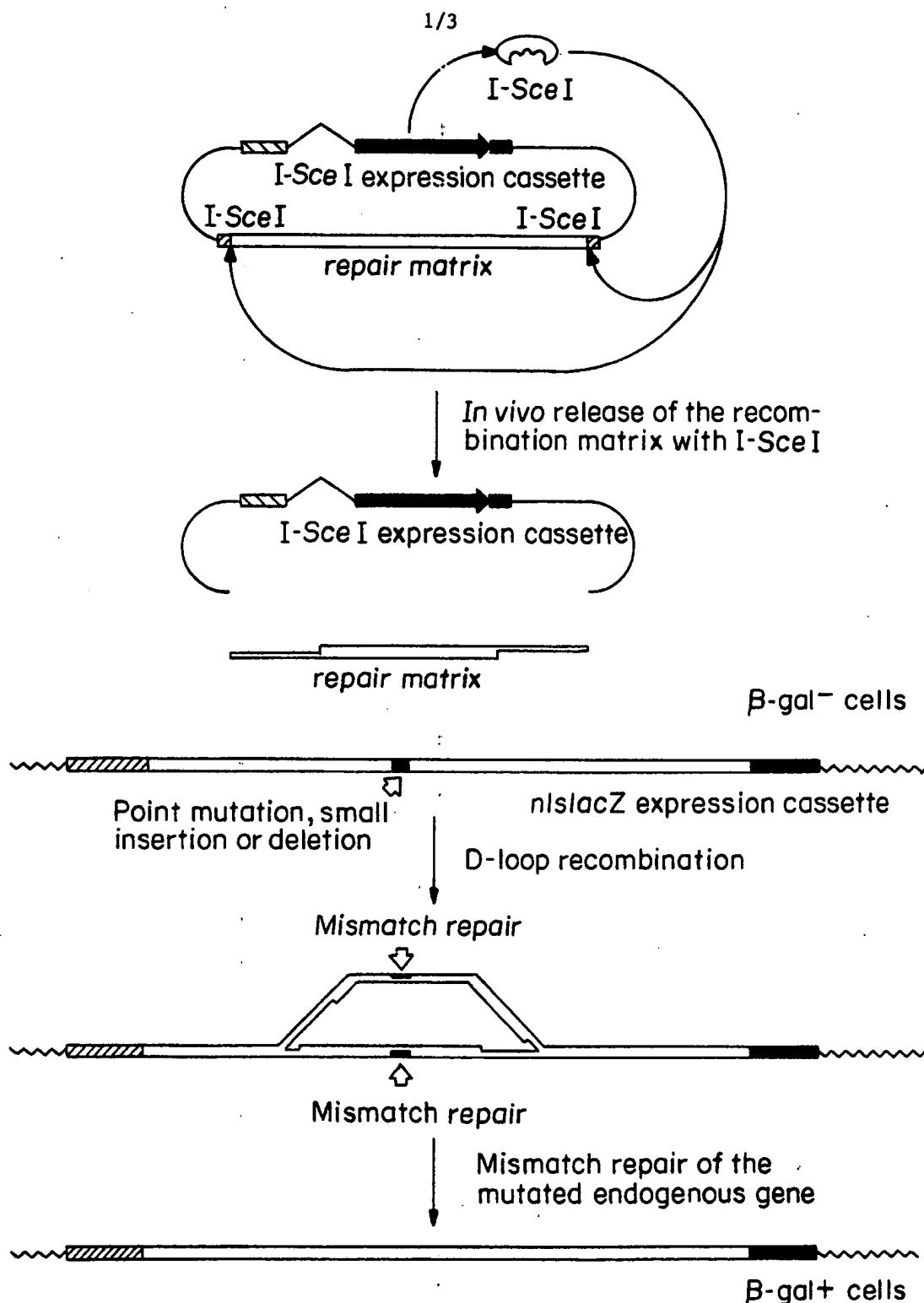


FIG. 1

2/3

% age of $\beta$ -gal expressing cells		No 1-SceI site	2 1-SceI sites	1 1-SceI sites	linear fragment
4 bp deletion repair	I-SceI +	0	0.8-1.2	0.09	0.001
	I-SceI -	0	0	0	0.001
4 bp duplication repair	I-SceI +	0	0-1.6	0.2-0.3	0.007
	I-SceI -	0	0	0	0.008

FIG. 2

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Enzyme	Organism (strain)	Size	Site	Cleavage	Shape of Cleavage	GenBank Accession No.	Commercially Available
<b>ENZYMES ENCODED BY INDEPENDENT GENES</b>							
Endo.Sce	Saccharomyces cerevisiae (IAM 4274)	476	25	very frequent	4/3'OH	M63839	
HO	Saccharomyces cerevisiae	586	18	very rare	4/3'OH	M14678	
<b>ENZYMES ENCODED BY INTRONS</b>							
I-Ceu I	Chlamydomonas eugametos	218	20	very rare	4/3'OH	S15138	Yes
I-Chu I	Chlamydomonas humicola	218	20	very rare	4/3'OH	L06107	
I-Cre I	Chlamydomonas reinhardtii	163	24	very rare	4/3'OH	X01977	
I-Csm I	Chlamydomonas smithii	237				X55305	
I-Dir I	Didymium iridis (Pan 2)	244				X71792	
I-Dmo I	Desulfurococcus mobilis	194		rare	4/3'OH	P21505	
I-Hmu I	Bacteriophage SP01/B. subtilis	174				M37686	
I-Hmu II	Bacteriophage SP82/B. subtilis	187					
I-Ppo I	Physarum polycephalum (Carolina)	185	15	rare	4/3'OH	M38131	Yes
I-Sce I	Saccharomyces cerevisiae (IL8-8C/R53)	235	18	very rare	4/3'OH	P03882	Yes
I-Sce II	Saccharomyces cerevisiae (D273-10B)	316	15	frequent	4/3'OH	P03878	
I-Sce III	Saccharomyces cerevisiae (777-3A)	335	18	very rare	4/3'OH	P03877	
I-Sce IV	Saccharomyces cerevisiae (777-3A)	307		rare	4/3'OH		
I-Tev I	Bacteriophage T4/E. coli	245	39	?	2/3'OH	M12742	
I-Tev II	Bacteriophage T4/E. coli	258	25	?	2/3'OH		
I-Tev III	Bacteriophage RB3/E. coli	269			2/5P	X59078	
<b>INTERCALATING PROTEINS</b>							
PI-Mle I	Mycobacterium leprae	365				X73822	
PI-Mtu I	Mycobacterium tuberculosis	439				X58485	
PI-Psp I	Pyrococcus species (GB-D)	537				U00707	
PI-Tli I	Thermococcus litoralis	390	20	rare	4/3'OH	M74198	Yes
PI-Tli II	Thermococcus litoralis	541	12	very rare	4/3'OH	M74198	Yes
PI-Sce V	Saccharomyces cerevisiae (YFG 499)	454	12	very rare	4/3'OH	M21609	Yes

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 00/02949

<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC 7 C12N15/90 //C12N9/22		
According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) IPC 7 C12N		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96 14408 A (PASTEUR INSTITUT ;UNIV PARIS CURIE (FR)) 17 May 1996 (1996-05-17) figure 30	1-55
A	CHOULIK A ET AL: "THE YEAST I-SCE I MEGANUCLEASE INDUCES SITE-DIRECTED CHROMOSOMAL RECOMBINATION IN MAMMALIAN CELLS" COMPTES RENDUS DES SEANCES DE L'ACADEMIE DES SCIENCES. SERIE III: SCIENCES DE LA VIE, NL, ELSEVIER, AMSTERDAM, vol. 317, no. 11, 1 November 1994 (1994-11-01), pages 1013-1019, XP000571580 ISSN: 0764-4469 the whole document	1-55
<input type="checkbox"/> Further documents are listed in the continuation of box C.		<input checked="" type="checkbox"/> Patent family members are listed in annex.
<p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"8" document member of the same patent family</p>		
Date of the actual completion of the international search		Date of mailing of the international search report
31 May 2000		07/06/2000
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentstaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Gallili, I

**INTERNATIONAL SEARCH REPORT**

International application No.  
PCT/US 00/02949

**Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)**

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)**

This International Searching Authority found multiple inventions in this International application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
  
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3.  As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

**Remark on Protest**

The additional search fees were accompanied by the applicant's protest.  
 No protest accompanied the payment of additional search fees.

**FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210**

**Continuation of Box I.1**

Claims 26-33 are explicitly directed to a method of treatment of the human/animal body. Moreover, claims 1-25,34-55 include methods *in vivo* and can therefore be directed to a method of treatment as well. The search has been carried out and based on the alleged effects of the compound/composition.

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

Int'l. Application No

PCT/US 00/02949

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9614408	A 17-05-1996	US 5792632 A		11-08-1998
		CA 2203569 A		17-05-1996
		EP 0791058 A		27-08-1997
		JP 10508478 T		25-08-1998
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		US 5866361 A		02-02-1999